PATENT

Support for the amendment of claim 1 herein can be found, inter alia, at page 3, lines 18-28. Support for the amendment of claim 13 herein can be found, inter alia, at page 9, line 4 -page 12, line 2.

## **REMARKS**

## Rejections Under 35 U.S.C. § 102(a)

The Examiner rejects claims 1, 2, 6, and 11 as unpatentable under 35 U.S.C. § 102(a) due to being anticipated by each of Timmermann et al., Kidney Intl. 53:1455-60 (June 1998) ("Timmerman #1") and Timmerman et al., J. Molecular Med. 76:B30, Abst. P-109 (May 1998) ("Timmerman #2").

In response to this rejection, Applicant submits herewith a Declaration of Prior Invention Under 37 C.F.R. § 1.131, demonstrating that the invention(s) embodied in these claims occurred prior to the effective date of either of these references. Applicant therefore respectfully requests the withdrawal of this rejection.

## Rejections Under 35 U.S.C. § 102(b)

The Examiner rejects claims 13 and 14 as unpatentable under 35 U.S.C. § 102(b) due to being anticipated by Intl. Publ. No. WO 97/35973.

In response to this rejection, Applicant submits herewith a Declaration of Prior Invention Under 37 C.F.R. § 1.131, demonstrating that the invention(s) embodied in these claims occurred prior to the effective date of this reference. Applicant therefore respectfully requests the withdrawal of this rejection.

Next, the Examiner rejects claims 13-15 as unpatentable under 35 U.S.C. § 102(b) due to being anticipated by each of (1) GenBank Accession No. Y00106 (Sept. 12, 1993), and (2) Emorine et al., Proc. Natl. Acad. Sci. USA 84:6995-9 (Oct. 1987) ("Emorine"). Although not

PATENT

acceding to the Examiner's rejection, Applicant notes that claim 13 has been amended herein to recite the fact that the composition comprises at least one allele-specific oligonucleotide (ASO) that hybridizes to a  $\beta_2$ AR polynucleotide at a region containing the 5'LC polymorphic site, wherein the ASO is not less than 10 nucleotides in length and not more than 100 nucleotides in length. Because the polynucleotide sequences disclosed in GenBank Accession No. Y00106 and in the Emorine reference are well in excess of 100 nucleotides, neither of these references anticipates claim 13 as amended. As such, Applicant respectfully requests the withdrawal of this rejection.

## Rejections Under 35 U.S.C. § 103(a)

The Examiner rejects claims 1, 2, 6-8, 11, 13, and 15 under 35 U.S.C. § 103(a) due to being rendered obvious by Timmermann et al., Kidney Intl. 53:1455-60 (June 1998)

(Timmerman #1) in view of Green et al., Amer. J. Resp. Cell Mol. Biol. 13:25-33 (July 1995)

("Green") and the Emorine reference.

In response to this rejection, Applicant submits herewith a Declaration of Prior Invention Under 37 C.F.R. § 1.131, demonstrating that the invention(s) embodied in these claims occurred prior to the effective date of the Timmerman #1 reference. Applicant therefore respectfully requests the withdrawal of this rejection.

Next, the Examiner rejects claims 1, 2, 6-8, 11, 13, and 15 under 35 U.S.C. § 103(a) due to being rendered obvious by the Timmerman #2 reference in view of the Green reference and the Emorine reference.

In response to this rejection, Applicant submits herewith a Declaration of Prior Invention Under 37 C.F.R. § 1.131, demonstrating that the invention(s) embodied in these claims occurred prior to the effective date of the Timmerman #2. Applicant therefore respectfully requests the withdrawal of this rejection.

Next, the Examiner rejects claims 1, 2, 6-8, 11, 13, and 15 under 35 U.S.C. § 103(a) due to being rendered obvious by Timmermann et al., Human Mutation 11(4):343-4 (March 1998) ("Timmerman #3") in view of the Green reference and the Emorine reference.

In response to this rejection, Applicant submits herewith a Declaration of Prior Invention Under 37 C.F.R. § 1.131, demonstrating that the invention(s) embodied in these claims occurred prior to the effective date of the Timmerman #3 reference. Applicant therefore respectfully requests the withdrawal of this rejection.

Next, the Examiner rejects claims 1, 2, 4-6, 11, 13, and 14 under 35 U.S.C. § 103(a) due to being rendered obvious by the Timmermann #1 reference in view of United States Patent No. 5,817,477 ("'477 patent").

In response to this rejection, Applicant submits herewith a Declaration of Prior Invention Under 37 C.F.R. § 1.131, demonstrating that the invention(s) embodied in these claims occurred prior to the effective date of the Timmerman #1 reference. Applicant therefore respectfully requests the withdrawal of this rejection.

Next, the Examiner rejects claims 1, 2, 4-6, 11, 13, and 14 under 35 U.S.C. § 103(a) due to being rendered obvious by the Timmermann #2 reference in view of the '477 patent and the Emorine reference.

In response to this rejection, Applicant submits herewith a Declaration of Prior Invention Under 37 C.F.R. § 1.131, demonstrating that the invention(s) embodied in these claims occurred prior to the effective date of the Timmerman #2 reference. Applicant therefore respectfully requests the withdrawal of this rejection.

Next, the Examiner rejects claims 1, 2, 4-6, 11, 13, and 14 under 35 U.S.C. § 103(a) due to being rendered obvious by the Timmermann #3 reference in view of the '477 patent and the Emorine reference.

In response to this rejection, Applicant submits herewith a Declaration of Prior Invention Under 37 C.F.R. § 1.131, demonstrating that the invention(s) embodied in these claims occurred

PATENT

prior to the effective date of the Timmerman #3 reference. Applicant therefore respectfully requests the withdrawal of this rejection.

Next, the Examiner rejects claims 1-3, 6, and 11 under 35 U.S.C. § 103(a) due to being rendered obvious by the Timmermann #3 reference in view of Large et al., J. Clin. Invest.

100:3005-13 (Dec. 1997) ("Large") and the Emorine reference, and further in view of New England BioLabs Catalog, p. 38 (1995) ("NEB Catalog").

In response to this rejection, Applicant submits herewith a Declaration of Prior Invention Under 37 C.F.R. § 1.131, demonstrating that the invention(s) embodied in these claims occurred prior to the effective date of the Timmerman #3 reference. Applicant therefore respectfully requests the withdrawal of this rejection.

Next, the Examiner rejects claims 1-3, 6, and 11 under 35 U.S.C. § 103(a) due to being rendered obvious by the Timmermann #2 reference in view of the Large reference and the Emorine reference, and further in view of the NEB Catalog reference.

In response to this rejection, Applicant submits herewith a Declaration of Prior Invention Under 37 C.F.R. § 1.131, demonstrating that the invention(s) embodied in these claims occurred prior to the effective date of the Timmerman #2 reference. Applicant therefore respectfully requests the withdrawal of this rejection.

Finally, the Examiner rejects claims 1 and 11 under 35 U.S.C. § 103(a) as unpatentable due to being rendered obvious by the Emorine reference in view of the '477 patent and United States Patent No. 6,087,485 ("'485 patent").

More specifically, the Examiner asserts that:

Emorine teaches methods of sequencing the  $\beta$ 2-AR gene and teachings [sic] the resulting sequence of the  $\beta$ 2-AR gene including the leader cistron sequences. Emorine does not teach determining the sequence of both copies of the  $\beta$ 2-AR gene.

However, ... [the '485 patent] teaches methods for detecting sequencing genomic DNA and for determining the presence of mutations and polymorphisms in DNA. IN [sic] the method of ... [the '485 patent], genomic DNA is amplified by PCR, cycle sequencing is performed, and sequences are determined and

analyzed for the presence of heterozygous positions. The method of . . . [the '485 patent] results in the analysis of the sequence of both copies of a genomic DNA sequence. Furthermore, . . . [the '477 patent] teaches the importance of determining the sequence of adrenergic receptor genes and of identifying the sequence variations in the adrenergic receptor genes.

In view of the teachings of ... [the '485 patent and the '477 patent], it would have been obvious to one of ordinary skill in the art at the time the invention was made to have analyzed the  $\beta$ 2-AR gene using the cycle sequencing method of . . . [the '485 patent] in order to have provided an effective means for analyzing the  $\beta$ 2-AR gene for the presence of genetic variation. Such a method would have analyzed all positions of the  $\beta$ 2-AR gene including the -47 5' LC polymorphic site and would have necessarily identified the nucleotide pair present at the 5' LC polymorphic site.

PTO Paper No. 13, pp. 20-21 (internal citations omitted).

Applicant respectfully disagrees with the Examiner's analysis. In order to establish a prima facie case of obviousness, the Examiner must establish three facts. First, the Examiner must establish that there exists some suggestion or motivation, either in the references themselves or in the knowledge generally available to one of ordinary skill in the art, to modify the reference or to combine reference teachings. Second, the Examiner must establish that there exists a reasonable expectation of success. Finally, the Examiner must establish that the prior art reference (or references when combined) teaches or suggests all the claim limitations. The teaching or suggestion to make the claimed combination and the reasonable expectation of success must both be found in the prior art, and not based on applicant's disclosure. In re Vaeck, 947 F.2d 488, 20 USPQ2d 1438 (Fed. Cir. 1991).

Without regard to whether the first two Vaeck criteria have been established, Applicant notes that the references – taken either individually or together – do not teach or suggest all the claim limitations of claims 1 and 11. Applicant has discovered the existence of DNA sequence variation with respect to the region upstream of the human  $\beta_2AR$  gene. Applicant has discovered that this variation occurs at a position 47 base pairs upstream of the coding region of the  $\beta_2AR$  gene, which begins at nucleotide position 1588 of SEQ ID NO:1. Applicant has, throughout the specification, defined this position as the "5' leader cistron polymorphic site."

PATENT

However, while the cited references may, in combination, teach or suggest a method for looking for polymorphic sites in the  $\beta_2AR$  gene, these references do not – taken either individually or together – teach or suggest a claim directed to genotyping the  $\beta_2AR$  gene of an individual comprising determining the identity of the nucleotide pair at a specific polymorphic site, namely the 5' leader cistron polymorphic site in the two copies of the  $\beta_2AR$  gene present in the individual. As such, the Examiner has not established a prima facie case of obviousness of claims 1 and 11 as filed and the rejection should be withdrawn.

Notwithstanding the foregoing, Applicant has, in an effort to expedite prosecution, amended claim 1 to merely clarify that the claimed method is indeed directed to determining the identity of the nucleotide pair at a specific  $\beta_2AR$  polymorphic site, which is not taught or suggested by the prior art, by reciting in the claim the nucleotide pairs that the specification states exist at that site, i.e., (a) cytosine and cytosine; (b) cytosine and thymine; and (c) thymine and thymine.

PATENT

Applicant notes that this response is being filed after the shortened statutory deadline for responding, April 7, 2003, but on or prior to one month following this deadline. Therefore, Applicant respectfully requests a one-month extension of time under 37 C.F.R. § 1.136(a), thereby extending the response period to May 7, 2003. Applicant authorizes the Commissioner to deduct the requisite fee for this extension (\$110.00; see 37 C.F.R. § 1.17(a)(2)) from Deposit Account No. 50-1293.

Respectfully submitted,

Matthew M. Catlett

Registration No. 44,067

Genaissance Pharmaceuticals, Inc.

Five Science Park

New Haven, CT 06511

203.776.1450

203.492.4474 (fax)